=> d l1 L1 HAS NO ANSWERS L1 STR

$$\begin{array}{c|c} & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & & \\ & & \\ & & & \\ & &$$

G1 X,OH,H

Structure attributes must be viewed using STN Express query preparation.

=> d his

(FILE 'HOME' ENTERED AT 09:57:05 ON 16 JAN 2008)

FILE 'REGISTRY' ENTERED AT 09:57:53 ON 16 JAN 2008

L1 STRUCTURE UPLOADED
L2 QUE L1
L3 2 S L1
L4 40 S L1 FUL
L5 30 S L4 AND CAPLUS/LC
L6 10 S L4 NOT L5

FILE 'CAPLUS' ENTERED AT 10:00:51 ON 16 JAN 2008

18 S L4 L7 L8. 14 S L4/THU L9 15 S L7 AND (GAST?) L10 1 S L9 AND FUND? 1 S L7 AND (BLOAT? OR ACCOMMADAT? OR PYLORIC?) Lll L121 S L11 NOT L10 3 S L7 AND (DYSPEP? OR INDIGEST?) L13 L14 2 S L13 NOT (L10 OR L12) 1 S L7 AND SATIE? L15 L16 0 S L15 NOT (L10 OR L12) L17 14 S L7 NOT (L10 OR L12 OR L14)



10/509,335 Page 1

L3 ANSWER 1 OF 1 SCISEARCH COPYRIGHT (c) 2008 The Thomson Corporation on

STN

ACCESSION NUMBER: 2003:274371 SCISEARCH

THE GENUINE ARTICLE: 657TV

TITLE: Z-338. Treatment of non-ulcer dyspepsia.

AUTHOR: Sorbera L A (Reprint); Castaner J; Leeson P A

CORPORATE SOURCE: Prous Sci, POB 540, Barcelona 08080, Spain (Reprint);

Prous Sci, Barcelona 08080, Spain

COUNTRY OF AUTHOR: Spain

SOURCE: DRUGS OF THE FUTURE, (JAN 2003) Vol. 28

, No. 1, pp. 26-30.

ISSN: 0377-8282.

PUBLISHER: PROUS SCIENCE, SA, PO BOX 540, PROVENZA 388, 08025

BARCELONA, SPAIN.

DOCUMENT TYPE: Article; Journal

LANGUAGE: English

REFERENCE COUNT: 30

ENTRY DATE: Entered STN: 11 Apr 2003

Last Updated on STN: 11 Apr 2003

ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS

Research efforts focusing on discovering gastroprokinetic agents with mechanisms of action different from available compounds have identified Z-338, a 2-(acylamino)thiazole-4-carboxamide derivative, as having potent gastroprokinetic activity and an excellent safety profile. Results from preclinical studies demonstrated that Z-338 enhances spontaneous contractions and electrically stimulated excitatory junction potentials and acetylcholine release, possibly through inhibition of muscarinic M-1 and M-2 autoreceptors and possibly an M5-like receptor. Z-338 has been shown to be safe in phase I trials involving healthy volunteers in Europe and Japan and in an early phase II trial conducted in patients with functional dyspepsia. Z-338 continues to undergo phase II development.

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L14 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2003:287963 CAPLUS <<LOGINID::20080116>>

DOCUMENT NUMBER: 139:30058

TITLE: Z-338: treatment of non-ulcer dyspepsia
AUTHOR(S): Sorbera, L. A.; Castaner, J.; Leeson, P. A.

CORPORATE SOURCE: Prous Science, Barcelona, 08080, Spain SOURCE: Drugs of the Future (2003), 28(1), 26-30

CODEN: DRFUD4; ISSN: 0377-8282

PUBLISHER: Prous Science

DOCUMENT TYPE: Journal; General Review

LANGUAGE: English

AB A review. Research efforts focusing on discovering gastroprokinetic agents with mechanisms of action different from available compds. have identified Z-338, a 2-(acylamino)thiazole-4-carboxamide derivative, as having potent gastroprokinetic activity and an excellent safety profile. Results from preclin. studies demonstrated that Z-338 enhances spontaneous contractions and elec. stimulated excitatory junction potentials and acetylcholine release, possibly through inhibition of muscarinic M1 and M2 autoreceptors and possibly an M5-like receptor. Z-338 has been shown to be safe in phase I trials involving healthy volunteers in Europe and Japan and in an early phase II trial conducted in patients with functional dyspepsia. Z-338 continues to undergo phase II development.

IT $\frac{185104-11-4P}{403651-06-9P}$ Z-338 $\frac{185106-16-5P}{403651-06-9P}$ $\frac{211999-70-1P}{403651-06-9P}$

RL: ADV (Adverse effect, including toxicity); PAC (Pharmacological activity); PKT (Pharmacokinetics); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prokinetic thiazolecarboxamide derivative Z-338 synthesis, pharmacokinetics, and pharmacol. activity)

RN 185104-11-4 CAPLUS

CN 4-Thiazolecarboxamide, N-[2-[bis(1-methylethyl)amino]ethyl]-2-[(2-hydroxy-4,5-dimethoxybenzoyl)amino]-, monohydrochloride (9CI) (CA INDEX NAME)

$$(i-Pr)_2N-CH_2-CH_2-NH-C NH-C NH-C OMe$$

HC1

RN 185106-16-5 CAPLUS

CN 4-Thiazolecarboxamide, N-[2-[bis(1-methylethyl)amino]ethyl]-2-[(2-hydroxy-4,5-dimethoxybenzoyl)amino]- (CA INDEX NAME)

$$(i-Pr)_2N-CH_2-CH_2-NH-C$$

$$NH-C$$

$$NH-C$$

$$NH-C$$

$$OMe$$

$$OMe$$

$$OMe$$

RN 211999-70-1 CAPLUS

CN 4-Thiazolecarboxamide, N-[2-[bis(1-methylethyl)amino]ethyl]-2-[(2-hydroxy-4,5-dimethoxybenzoyl)amino]-, (2Z)-2-butenedioate (1:1) (salt) (9CI) (CA INDEX NAME)

CM. 1

CRN 185106-16-5 CMF C21 H30 N4 O5 S

$$(i-Pr)_2N-CH_2-CH_2-NH-C NH-C NH-C OMe$$

CM 2

CRN 110-16-7 CMF C4 H4 O4

Double bond geometry as shown.

RN 403651-06-9 CAPLUS

CN 4-Thiazolecarboxamide, N-[2-[bis(1-methylethyl)amino]ethyl]-2-[(2-hydroxy-4,5-dimethoxybenzoyl)amino]-, trihydrate (9CI) (CA INDEX NAME)

$$(i-Pr)_2N-CH_2-CH_2-NH-C NH-C NH-C OMe$$

●3 H₂O

REFERENCE COUNT:

THERE ARE 27 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT